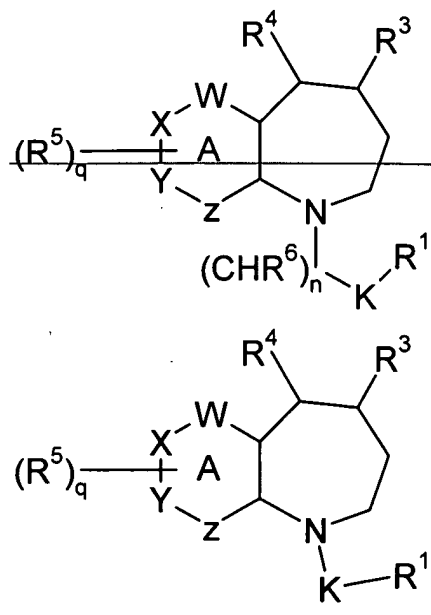


Amendments to the Claims

1. (currently amended) A compound of a formula below



wherein

~~n is 0, 1, 2, or 3;~~

q is 0, 1, or 2;

W, X, Y and Z are each independently CH, C, N, or S, ~~or O~~ with appropriate single or double bonds and/or hydrogen atoms to complete valency requirements providing Ring A as a five or six member ring, wherein one of W, X, Y or Z may be absent, selected from pyridine, thiophene, or pyrazole;

K is a bond or C=O;

p is 0, 1 or 2;

R<sup>1</sup> is selected from a group consisting of hydroxy, hydrogen, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>3</sub>-C<sub>8</sub> cycloalkyl, aryl, C<sub>2</sub>-C<sub>6</sub> alkylalcohol, -OC<sub>1</sub>-C<sub>6</sub> alkyl, -O-aryl, -OC<sub>3</sub>-C<sub>8</sub> cycloalkyl, and NR<sup>7</sup>R<sup>8</sup> wherein each cycloalkyl and aryl is optionally substituted with C<sub>0</sub>-C<sub>6</sub> alkylCOOR<sup>11</sup>;

R<sup>3</sup> is hydrogen or C<sub>1</sub>-C<sub>6</sub> alkyl;

R<sup>4</sup> is a group represented by the formula -NR<sup>9</sup>R<sup>10</sup>;

R<sup>5</sup> is selected from the group consisting of hydrogen, halogen, hydroxy, C<sub>1</sub>-C<sub>6</sub> alkyl, OC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> haloalkyl, and NR<sup>7</sup>R<sup>8</sup>, ~~and~~ -CN;

R<sup>7</sup> and R<sup>8</sup> are ~~independently selected~~ hydrogen or C<sub>1</sub>-C<sub>6</sub> alkyl;

$R^9$  is selected from:  $COR^7$ ,  $CO_2R^7$ ,  $C_0-C_3$  alkylCONR<sup>7</sup>R<sup>8</sup>,  $C_0-C_3$  alkylS(O)<sub>p</sub>NR<sup>7</sup>R<sup>8</sup>, or  $C_0-C_3$  alkylS(O)<sub>p</sub>R<sup>7</sup>, or tetrazole optionally substituted with one or two  $C_1-C_6$  alkyl groups;

$R^{10}$  is  $C_1-C_6$  alkylaryl optionally substituted with 1-3 groups independently selected from  $C_1-C_6$  alkyl,  $C_1-C_6$  haloalkyl, halogen, or cyano;

$R^{11}$  is hydrogen or  $C_1-C_6$  alkyl;

or a pharmaceutically acceptable salt thereof.

2. (currently amended) A compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein n is 0, and wherein  $R^1$  is selected from  $C_1-C_6$  alkyl, or  $-OC_1-C_6$  alkyl.

3. (previously presented) A compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein  $R^9$  is tetrazole optionally substituted with one or two  $C_1-C_6$  alkyl groups.

4. (canceled)

5. (currently amended) A compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein ~~n and~~ q are independently 0 or 1.

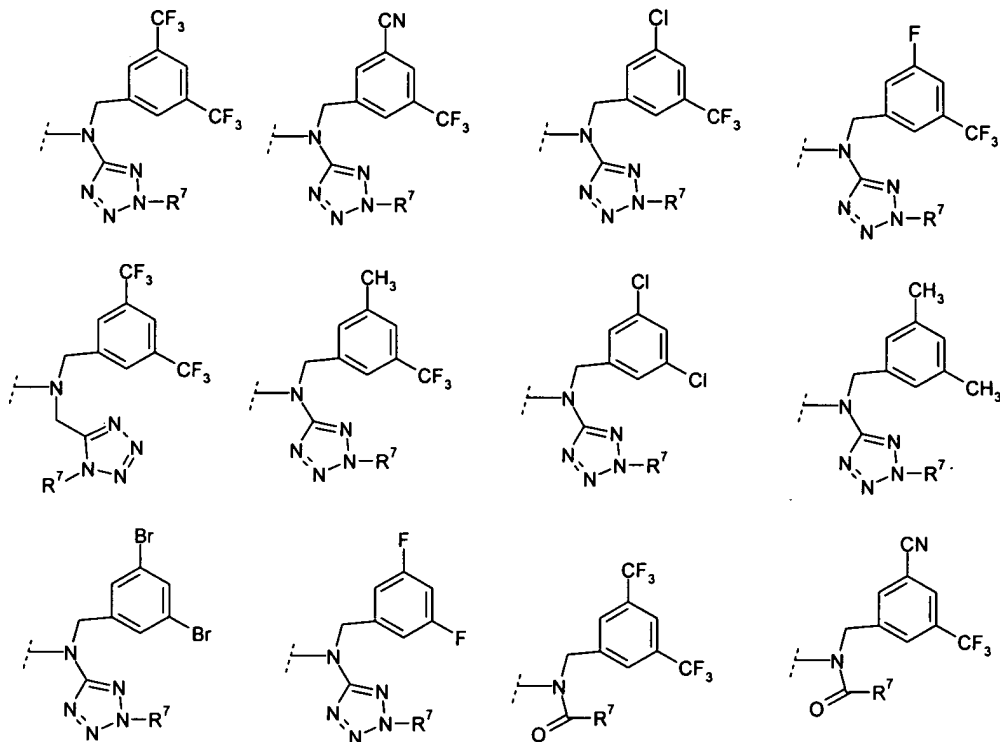
6. (previously presented) A compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein the A ring is pyridine or thiophene.

7. (previously presented) A compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein the A ring is pyridine.

8. (previously presented) A compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein the A ring is thiophene.

9. (canceled)

10. (previously presented) A compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein and  $R^4$  is selected from the group consisting of:



wherein R<sup>7</sup> is C<sub>1</sub>-C<sub>6</sub> alkyl.

11. (previously presented) A compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein R<sup>9</sup> is COOR<sup>7</sup>.

12. (previously presented) A compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein R<sup>9</sup> is CONR<sup>7</sup>R<sup>8</sup>.

13. (previously presented) A compound according to Claim 1, or a pharmaceutically acceptable salt thereof, wherein R<sup>9</sup> is S(O)<sub>2</sub>NR<sup>7</sup>R<sup>8</sup>.

14. (currently amended) A compound selected from the group consisting of:  
 5-[Acetyl-(3,5-bis-trifluoromethyl-benzyl)-amino]-2,3,4,5-tetrahydro-thieno[3,4-b]azepine-1-carboxylic acid isopropyl ester,  
 8-[Acetyl-(3,5-bis-trifluoromethyl-benzyl)-amino]-3-methyl-5,6,7,8-tetrahydro-thieno[3,2-b]azepine-4-carboxylic acid isopropyl ester  
 8-[Acetyl-(3,5-bis-trifluoromethyl-benzyl)-amino]-2-bromo-5,6,7,8-tetrahydro-thieno[3,2-b]azepine-4-carboxylic acid isopropyl ester,

5-[Acetyl-(3,5-bis-trifluoromethyl-benzyl)-amino]-5,6,7,8-tetrahydro-pyrido[2,3-b]azepine-9-carboxylic acid isopropyl ester,  
5-[Acetyl-(3,5-bis-trifluoromethyl-benzyl)-amino]-2,3,4,5-tetrahydro-pyrido[3,4-b]azepine-1-carboxylic acid isopropyl ester,  
5-[Acetyl-(3,5-bis-trifluoromethyl-benzyl)-amino]-2,3,4,5-tetrahydro-pyrido[4,3-b]azepine-1-carboxylic acid isopropyl ester,  
9-[Acetyl-(3,5-bis-trifluoromethyl-benzyl)-amino]-6,7,8,9-tetrahydro-pyrido[3,2-b]azepine-5-carboxylic acid isopropyl ester,  
9-[Acetyl-(3,5-bis-trifluoromethyl-benzyl)-amino]-2-trifluoromethyl-6,7,8,9-tetrahydro-pyrido[3,2-b]azepine-5-carboxylic acid isopropyl ester,  
9-[Acetyl-(3,5-bis-trifluoromethyl-benzyl)-amino]-3-trifluoromethyl-6,7,8,9-tetrahydro-pyrido[3,2-b]azepine-5-carboxylic acid isopropyl ester,  
5-[Acetyl-(3,5-bis-trifluoromethyl-benzyl)-amino]-2,3,4,5-tetrahydro-thieno[3,4-b]azepine-1-carboxylic acid isopropyl ester,  
8-[Acetyl-(3,5-bis-trifluoromethyl-benzyl)-amino]-3-methyl-5,6,7,8-tetrahydro-thieno[3,2-b]azepine-4-carboxylic acid isopropyl ester,  
4-[Acetyl-(3,5-bis-trifluoromethyl-benzyl)-amino]-1-methyl-4,5,6,7-tetrahydro-1H-1,2,8-triaza-azulene-8-carboxylic acid isopropyl ester,  
9-[acetyl-(3,5-bis-trifluoromethylbenzyl)amino]-2-chloro-6,7,8,9-tetrahydro-pyrido[3,2-b]azepine-5-carboxylic acid isopropyl ester,  
9-[acetyl-(3,5-bis-trifluoromethylbenzyl)amino]-2-methoxy-6,7,8,9-tetrahydro-pyrido[3,2-b]azepine-5-carboxylic acid isopropyl ester,  
9-[acetyl-(3,5-bis-trifluoromethylbenzyl)amino]-2-bromo-6,7,8,9-tetrahydro-pyrido[3,2-b]azepine-5-carboxylic acid isopropyl ester,  
9-[Acetyl-(3,5-bis-trifluoromethylbenzyl)amino]-2-dimethylamino-6,7,8,9-tetrahydro-pyrido[3,2-b]azepine-5-carboxylic acid isopropyl ester,  
9-[Acetyl-(3,5-bis-trifluoromethylbenzyl)amino]-2-methyl-6,7,8,9-tetrahydro-pyrido[3,2-b]azepine-5-carboxylic acid isopropyl ester,  
9-[Acetyl-(3,5-bis-trifluoromethylbenzyl)amino]-2-cyano-6,7,8,9-tetrahydro-pyrido[3,2-b]azepine-5-carboxylic acid isopropyl ester,  
9-[Acetyl-(3,5-bis-trifluoromethylbenzyl)amino]-3-chloro-2-methoxy-6,7,8,9-tetrahydro-pyrido[3,2-b]azepine-5-carboxylic acid isopropyl ester,  
9-[Acetyl-(3,5-bis-trifluoromethylbenzyl)amino]-3-chloro-2-ethoxy-6,7,8,9-tetrahydro-pyrido[3,2-b]azepine-5-carboxylic acid isopropyl ester,

9-[Acetyl-(3,5-bis-trifluoromethyl-benzyl)amino]-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-pyrido[3,2-*b*]azepine-5-carboxylic acid isopropyl ester,  
9-[Acetyl-(3,5-bis-trifluoromethyl-benzyl)amino]-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-pyrido[3,2-*b*]azepine-5-carboxylic acid *tert*-butyl ester,  
9-[(3,5-Bis-trifluoromethyl-benzyl)-2-methyl-2*H*-tetrazol-5-yl)-amino]-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-pyrido[3,2-*b*]azepine-5-carboxylic acid isopropyl ester,  
9-[(3,5-Bis-trifluoromethyl-benzyl)-2-methyl-2*H*-tetrazol-5-yl)-amino]-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-pyrido[3,2-*b*]azepine-5-carboxylic acid *tert*-butyl ester,  
(3,5-Bis-trifluoromethyl-benzyl)-(5-cyclopentylmethyl-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-5*H*-pyrido[3,2-*b*]azepin-9-yl)-(2-methyl-2*H*-tetrazol-5-yl)-amine,  
(3,5-Bis-trifluoromethyl-benzyl)-(5-cyclopropylmethyl-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-5*H*-pyrido[3,2-*b*]azepin-9-yl)-(2-methyl-2*H*-tetrazol-5-yl)-amine,  
(3,5-Bis-trifluoromethyl-benzyl)-(2-methyl-5-pyridin-3-ylmethyl-3-trifluoromethyl-6,7,8,9-tetrahydro-5*H*-pyrido[3,2-*b*]azepin-9-yl)-(2-methyl-2*H*-tetrazol-5-yl)-amine,  
(3,5-Bis-trifluoromethyl-benzyl)-(2-methyl-5-pyridin-4-ylmethyl-3-trifluoromethyl-6,7,8,9-tetrahydro-5*H*-pyrido[3,2-*b*]azepin-9-yl)-(2-methyl-2*H*-tetrazol-5-yl)-amine,  
3-{9-[(3,5-Bis-trifluoromethyl-benzyl)-(2-methyl-2*H*-tetrazol-5-yl)-amino]-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-pyrido[3,2-*b*]azepin-5-ylmethyl}-benzoic acid,  
4-{9-[(3,5-Bis-trifluoromethyl-benzyl)-(2-methyl-2*H*-tetrazol-5-yl)-amino]-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-pyrido[3,2-*b*]azepin-5-ylmethyl}-benzoic acid,  
5-{9-[(3,5-Bis-trifluoromethyl-benzyl)-(2-methyl-2*H*-tetrazol-5-yl)-amino]-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-pyrido[3,2-*b*]azepin-5-yl}-3,3-dimethyl-pentanoic acid,  
(4-{9-[(3,5-Bis-trifluoromethyl-benzyl)-(2-methyl-2*H*-tetrazol-5-yl)-amino]-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-pyrido[3,2-*b*]azepin-5-ylmethyl}-cyclohexyl)-acetic acid,  
(3,5-Bis-trifluoromethyl-benzyl)-(5-ethyl-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-5*H*-pyrido[3,2-*b*]azepin-9-yl)-(2-methyl-2*H*-tetrazol-5-yl)-amine,  
5-{9-[(3,5-Bis-trifluoromethyl-benzyl)-(2-methyl-2*H*-tetrazol-5-yl)-amino]-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-pyrido[3,2-*b*]azepin-5-ylmethyl}-thiophene-2-carboxylic acid,  
2-{9-[(3,5-Bis-trifluoromethyl-benzyl)-(2-methyl-2*H*-tetrazol-5-yl)-amino]-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-pyrido[3,2-*b*]azepin-5-yl}-ethanol,  
(5-Benzyl-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-5*H*-pyrido[3,2-*b*]azepin-9-yl)-(3,5-bis-trifluoromethyl-benzyl)-(2-methyl-2*H*-tetrazol-5-yl)-amine,  
(3,5-Bis-trifluoromethyl-benzyl)-(2-methyl-2*H*-tetrazol-5-yl)-(2-methyl-5-thiazol-2-ylmethyl-3-trifluoromethyl-6,7,8,9-tetrahydro-5*H*-pyrido[3,2-*b*]azepin-9-yl)-amine,

9-[(3,5-Bis-trifluoromethyl-benzyl)-(2-methyl-2H-tetrazol-5-yl)-amino]-2-methyl-3-trifluoromethyl-6,7,8,9-tetrahydro-pyrido[3,2-b]azepine-5-carboxylic acid tetrahydro-furan-3-yl ester,

~~(3,5-Bis-trifluoromethyl-benzyl)-(2-methyl-5-pyridin-4-ylmethyl-3-trifluoromethyl-6,7,8,9-tetrahydro-5H-pyrido[3,2-b]azepin-9-yl)-carbamic acid methyl ester,~~

~~N-(3,5-Bis-trifluoromethyl-benzyl)-N-(2-methyl-5-pyridin-4-ylmethyl-3-trifluoromethyl-6,7,8,9-tetrahydro-5H-pyrido[3,2-b]azepin-9-yl)-acetamide~~

or a pharmaceutically acceptable salt thereof.

15-16. (canceled)

17. (previously presented) A method of treating atherosclerosis comprising administering a compound of formula I according to claim 1, or a pharmaceutically acceptable salt thereof to a patient.

18-20. (canceled)

21. (previously presented) A pharmaceutical composition comprising a compound according to Claim 1, or a pharmaceutically acceptable salt thereof, and at least one of a carrier, diluent and excipient.

22-28. (canceled)

29. (previously presented) A method of raising plasma HDL-cholesterol in a mammal comprising administering a therapeutically effective dose of a compound according to claim 1, or a pharmaceutically acceptable salt thereof to said mammal.

30. (previously presented) A pharmaceutical composition of claim 21 comprising one or more cardio protective agents selected from the group consisting of: statins, leptin, and lipid regulating agents.

31. (new) A method of treating a patient in need thereof for dyslipidemia comprising administering a compound of formula I according to claim 1, or a pharmaceutically acceptable salt thereof to the patient.